

THE CLAIMS

The listing of claims will replace all prior versions of claims in the application.

Listing of Claims:

1. (Currently amended) A method ~~to protect cells in a lipid bilayer membrane of~~ treating cell oxidative damage in humans, comprising administering a formulation comprising:
Vitamin E as d- α -tocopherol;
Vitamin E as dl- α -tocopheryl;
Vitamin E mixed tocopherols; and
tocotrienols in the forms comprising inseparable tocopherols.
2. (Original) The method of claim 1 wherein said tocotrienols are in the forms α , γ , β , and δ , and said inseparable tocopherols are in the forms of α , γ , β , and δ , said tocotrienols and said tocopherols being from rice, whereby said formulation is beneficial for antioxidant protection of cells in the human body containing a lipid layer membrane.
3. (Original) The method of claim 1 wherein said tocotrienols are in the forms α , γ , β , and δ , and said inseparable tocopherols are in the forms of α , γ , β , and δ , said tocotrienols and said tocopherols being from palm, whereby said formulation is beneficial for antioxidant protection of cells in the human body containing a lipid layer membrane.
4. (Original) The method of claim 1 wherein said Vitamin E mixed tocopherols are in the forms α , γ , β , and δ and are a blend of synthetic and natural sources of Vitamin E.
5. (Original) The method of claim 1 wherein said Vitamin E dl- α -tocopheryl is present at about 90 weight % of said active ingredients.
6. (Original) The method of claim 1 wherein said Vitamin E mixed tocopherols are present at about 5 weight % of said active ingredients.

7. (Original) The method of claim 1 wherein said tocotrienols from natural sources are present at about 5 weight % of said active ingredients.

8. (Currently amended) A method ~~to protect cells in a lipid bilayer membrane of~~ treating cell oxidative damage in humans, comprising administering a formulation comprising:

Vitamin E selected from at least one of the ester group consisting of:

dl- α -tocopheryl acetate; and

dl- α -tocopheryl succinate;

Vitamin E as d- α -tocopherol;

Vitamin E mixed tocopherols in the forms α , β , γ , and δ ;

tocotrienols in the forms α , β , γ , and δ .

9. (Original) The method of claim 8 wherein said Vitamin E as dl- α -tocopheryl ester, said Vitamin E as d- α -tocopherol, and said Vitamin E mixed tocopherols in the forms α , β , γ , and δ is a blend of synthetic and natural sources of Vitamin E, and said tocotrienols are from a natural source.

10. (Original) The method of claim 8 wherein said Vitamin E as dl- α -tocopheryl ester is present at from 5 mg to 400 mg.

11. (Original) The method of claim 8 wherein said Vitamin E as d- α -tocopherol is present at from 5 mg to 400 mg.

12. (Original) The method of claim 8 wherein said Vitamin E as mixed tocopherols is present at from 5 mg to 200 mg.

13. (Original) The method of claim 8 wherein said Vitamin E as mixed tocotrienols in the forms α , β , γ , and δ is present at from 5 mg to 50 mg with variable composition of isomers:

α tocotrienol at 1 to 30%;

β tocotrienol at 0.1 to 30%;

γ tocotrienol at 1 to 30%; and
 δ tocotrienol at 0.1 to 30%.

14. (Original) The method of claim 13 comprising: inseparable variable content of carotenoids comprising:

alpha carotene;
beta carotene;
gamma carotene;
lycopene; and
phytosterols and squalene.

15. (Original) The method of claim 8 comprising:
a marker selected from at least one of the group consisting of:

coenzyme Q10;
rosemary oil;
green tea;
 α lipoic acid;
lycopene;
grape seed extract;
pine bark extract;
vitamin C;
natural beta carotene;
synthetic beta carotene;
 γ -oryzanol;
selenium; and
lutein.

16. (Currently amended) A method ~~to protect cells in a lipid bilayer membrane of~~
treating cell oxidative damage in humans, comprising administering a formulation comprising:

Vitamin E selected from at least one of the ester group consisting of:
dl- α -tocopheryl acetate; and

dl- α -tocopheryl succinate;
Vitamin E as d- α -tocopherol;
Vitamin E mixed tocopherols in the forms α , β , γ , and δ ; and tocotrienols in the forms α , β , γ , and δ .

17. (Original) The method of claim 16 wherein said Vitamin E as dl- α -tocopheryl ester, said Vitamin E as d- α -tocopherol, and said Vitamin E mixed tocopherols in the forms α , β , γ , and δ is a blend of synthetic and natural sources of Vitamin E, and said tocotrienols are from a natural source.

18. (Original) The method of claim 16 wherein said Vitamin E as dl- α -tocopheryl ester is present at from 5 mg to 2000 mg.

19. (Original) The method of claim 16 wherein said Vitamin E as d- α -tocopherol is present at from 5 mg to 2000 mg.

20. (Original) The method of claim 16 wherein said Vitamin E as mixed tocopherols is present at from 5 mg to 2000 mg.

21. (Original) The method of claim 16 wherein said Vitamin E as mixed tocotrienols in the forms α , β , γ , and δ is present at from 5 mg to 500 mg with variable composition of isomers:

α tocotrienol at 1 to 30%;
 β tocotrienol at 0.1 to 30%;
 γ tocotrienol at 1 to 30%; and
 δ tocotrienol at 0.1 to 30%.

22. (Original) The method of claim 16 wherein said Vitamin E as dl- α -tocopheryl ester, said Vitamin E as d- α -tocopherol, and said Vitamin E mixed tocopherols in the forms α , β , γ , and δ is a blend of synthetic and natural sources of Vitamin E, and said tocotrienols are from a natural source.

23. (Original) The method of claim 16 wherein said formulation is formed in a soft gel capsule further comprising :

gelatin;
glycerin; and
water for said soft gelatin formulation.

24. (Original) The method of claim 16 comprising: a marker selected from at least one of the group consisting of:

coenzyme Q10;
rosemary oil;
green tea;
 α lipoic acid;
lycopene;
grape seed extract;
pine bark extract;
vitamin C;
natural beta carotene;
synthetic beta carotene;
 γ -oryzanol;
selenium; and
lutein.

25. (Canceled)

26. (New) A method of protecting against cell oxidative damage in humans, comprising administering a formulation comprising:

Vitamin E as d- α -tocopherol;
Vitamin E as dl- α -tocopheryl;
Vitamin E mixed tocopherols; and
tocotrienols in the forms comprising inseparable tocopherols.

27. (New) The method of claim 26 wherein said tocotrienols are in the forms α , γ , β , and δ , and said inseparable tocopherols are in the forms of α , γ , β , and δ , said tocotrienols and said tocopherols being from rice, whereby said formulation is beneficial for antioxidant protection of cells in the human body containing a lipid layer membrane.

28. (New) The method of claim 26 wherein said tocotrienols are in the forms α , γ , β , and δ , and said inseparable tocopherols are in the forms of α , γ , β , and δ , said tocotrienols and said tocopherols being from palm, whereby said formulation is beneficial for antioxidant protection of cells in the human body containing a lipid layer membrane.

29. (New) The method of claim 26 wherein said Vitamin E mixed tocopherols are in the forms α , γ , β , and δ and are a blend of synthetic and natural sources of Vitamin E.

30. (New) The method of claim 26 wherein said Vitamin E dl- α -tocopheryl is present at about 90 weight % of said active ingredients.

31. (New) The method of claim 26 wherein said Vitamin E mixed tocopherols are present at about 5 weight % of said active ingredients.

32. (New) The method of claim 26 wherein said tocotrienols from natural sources are present at about 5 weight % of said active ingredients.

33. (New) A method of protecting against cell oxidative damage in humans, comprising administering a formulation comprising:

Vitamin E selected from at least one of the ester group consisting of:

dl- α -tocopheryl acetate; and

dl- α -tocopheryl succinate;

Vitamin E as d- α -tocopherol;

Vitamin E mixed tocopherols in the forms α , β , γ , and δ ;

tocotrienols in the forms α , β , γ , and δ .

34. (New) The method of claim 33 wherein said Vitamin E as dl- α -tocopheryl ester, said Vitamin E as d- α -tocopherol, and said Vitamin E mixed tocopherols in the forms α , β , γ , and δ is a blend of synthetic and natural sources of Vitamin E, and said tocotrienols are from a natural source.

35. (New) The method of claim 33 wherein said Vitamin E as dl- α -tocopheryl ester is present at from 5 mg to 400 mg.

36. (New) The method of claim 33 wherein said Vitamin E as d- α -tocopherol is present at from 5 mg to 400 mg.

37. (New) The method of claim 33 wherein said Vitamin E as mixed tocopherols is present at from 5 mg to 200 mg.

38. (New) The method of claim 33 wherein said Vitamin E as mixed tocotrienols in the forms α , β , γ , and δ is present at from 5 mg to 50 mg with variable composition of isomers:

- α tocotrienol at 1 to 30%;
- β tocotrienol at 0.1 to 30%;
- γ tocotrienol at 1 to 30%; and
- δ tocotrienol at 0.1 to 30%.

39. (New) The method of claim 38 comprising: inseparable variable content of carotenoids comprising:

- alpha carotene;
- beta carotene;
- gamma carotene;
- lycopene; and
- phytosterols and squalene.

40. (New) The method of claim 33 comprising:
a marker selected from at least one of the group consisting of:
coenzyme Q10;
rosemary oil;
green tea;
 α lipoic acid;
lycopene;
grape seed extract;
pine bark extract;
vitamin C;
natural beta carotene;
synthetic beta carotene;
 γ -oryzanol;
selenium; and
lutein.

41. (New) A method of protecting against cell oxidative damage in humans,
comprising administering a formulation comprising:

Vitamin E selected from at least one of the ester group consisting of:

dl- α -tocopheryl acetate; and

dl- α -tocopheryl succinate;

Vitamin E as d- α -tocopherol;

Vitamin E mixed tocopherols in the forms α , β , γ , and δ ; and tocotrienols in the forms α , β , γ , and δ .

42. (New) The method of claim 41 wherein said Vitamin E as dl- α -tocopheryl ester, said Vitamin E as d- α -tocopherol, and said Vitamin E mixed tocopherols in the forms α , β , γ , and δ is a blend of synthetic and natural sources of Vitamin E, and said tocotrienols are from a natural source.

43. (New) The method of claim 41 wherein said Vitamin E as dl- α -tocopheryl ester is present at from 5 mg to 2000 mg.

44. (New) The method of claim 41 wherein said Vitamin E as d- α -tocopherol is present at from 5 mg to 2000 mg.

45. (New) The method of claim 41 wherein said Vitamin E as mixed tocopherols is present at from 5 mg to 2000 mg.

46. (New) The method of claim 41 wherein said Vitamin E as mixed tocotrienols in the forms α , β , γ , and δ is present at from 5 mg to 500 mg with variable composition of isomers:

α tocotrienol at 1 to 30%;

β tocotrienol at 0.1 to 30%;

γ tocotrienol at 1 to 30%; and

δ tocotrienol at 0.1 to 30%.

47. (New) The method of claim 41 wherein said Vitamin E as dl- α -tocopheryl ester, said Vitamin E as d- α -tocopherol, and said Vitamin E mixed tocopherols in the forms α , β , γ , and δ is a blend of synthetic and natural sources of Vitamin E, and said tocotrienols are from a natural source.

48. (New) The method of claim 41 wherein said formulation is formed in a soft gel capsule further comprising :

gelatin;

glycerin; and

water for said soft gelatin formulation.

49. (New) The method of claim 41 comprising: a marker selected from at least one of the group consisting of:

coenzyme Q10;
rosemary oil;
green tea;
 α lipoic acid;
lycopene;
grape seed extract;
pine bark extract;
vitamin C;
natural beta carotene;
synthetic beta carotene;
 γ -oryzanol;
selenium; and
lutein.